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* * * * * * * * *
                     Welcome to STN International
                 Web Page for STN Seminar Schedule - N. America
NEWS
      1
                 CHEMLIST enhanced with New Zealand Inventory of Chemicals
NEWS
         JAN 08
NEWS
      3
         JAN 16
                 CA/CAplus Company Name Thesaurus enhanced and reloaded
NEWS
     4
         JAN 16
                 IPC version 2007.01 thesaurus available on STN
NEWS
     5
         JAN 16
                 WPIDS/WPINDEX/WPIX enhanced with IPC 8 reclassification data
NEWS
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         JAN 22
                 CA/CAplus updated with revised CAS roles
NEWS
     7
         JAN 22
                 CA/CAplus enhanced with patent applications from India
NEWS
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         JAN 29
                 PHAR reloaded with new search and display fields
NEWS 9
         JAN 29
                 CAS Registry Number crossover limit increased to 300,000 in
                 multiple databases
NEWS 10
         FEB 15
                 PATDPASPC enhanced with Drug Approval numbers
                 RUSSIAPAT enhanced with pre-1994 records
NEWS 11
        FEB 15
                 KOREAPAT enhanced with IPC 8 features and functionality
NEWS 12 FEB 23
NEWS 13 FEB 26 MEDLINE reloaded with enhancements
NEWS 14 FEB 26 EMBASE enhanced with Clinical Trial Number field
NEWS 15 FEB 26
                 TOXCENTER enhanced with reloaded MEDLINE
NEWS 16 FEB 26
                 IFICDB/IFIPAT/IFIUDB reloaded with enhancements
NEWS 17 FEB 26 CAS Registry Number crossover limit increased from 10,000
                 to 300,000 in multiple databases
NEWS 18 MAR 15
                 WPIDS/WPIX enhanced with new FRAGHITSTR display format
NEWS 19 MAR 16
                CASREACT coverage extended
NEWS 20 MAR 20 MARPAT now updated daily
NEWS 21 MAR 22 LWPI reloaded
NEWS 22 MAR 30 RDISCLOSURE reloaded with enhancements
NEWS 23 APR 02 JICST-EPLUS removed from database clusters and STN
NEWS 24 APR 30 GENBANK reloaded and enhanced with Genome Project ID field
NEWS 25 APR 30
                CHEMCATS enhanced with 1.2 million new records
NEWS 26 APR 30
                 CA/CAplus enhanced with 1870-1889 U.S. patent records
NEWS 27
        APR 30
                 INPADOC replaced by INPADOCDB on STN
NEWS 28
        MAY 01
                 New CAS web site launched
NEWS 29
         MAY 08
                 CA/CAplus Indian patent publication number format defined
NEWS 30
        MAY 14
                 RDISCLOSURE on STN Easy enhanced with new search and display
                 fields
             NOVEMBER 10 CURRENT WINDOWS VERSION IS V8.01c, CURRENT
NEWS EXPRESS
              MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
              AND CURRENT DISCOVER FILE IS DATED 25 SEPTEMBER 2006.
NEWS HOURS
              STN Operating Hours Plus Help Desk Availability
              Welcome Banner and News Items
NEWS LOGIN
```

Enter NEWS followed by the item number or name to see news on that specific topic.

For general information regarding STN implementation of IPC 8

NEWS IPC8

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FILE 'HOME' ENTERED AT 10:58:33 ON 18 MAY 2007

=>
Uploading
THIS COMMAND NOT AVAILABLE IN THE CURRENT FILE
Do you want to switch to the Registry File?
Choice (Y/n):
Switching to the Registry File...

Some commands only work in certain files. For example, the EXPAND command can only be used to look at the index in a file which has an index. Enter "HELP COMMANDS" at an arrow prompt (=>) for a list of commands which can be used in this file.

=> FILE REGISTRY

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 0.21 0.21

FULL ESTIMATED COST

FILE 'REGISTRY' ENTERED AT 10:58:47 ON 18 MAY 2007
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PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
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Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 17 MAY 2007 HIGHEST RN 935249-87-9 DICTIONARY FILE UPDATES: 17 MAY 2007 HIGHEST RN 935249-87-9

New CAS Information Use Policies, enter HELP USAGETERMS for details.

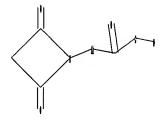
TSCA INFORMATION NOW CURRENT THROUGH December 2, 2006

Please note that search-term pricing does apply when conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/support/stngen/stndoc/properties.html

=>
Uploading C:\Program Files\Stnexp\Queries\10519835.str



chain nodes :

5 6 7 8 9 10 11

ring nodes : 1 2 3 4

chain bonds :

1-6 3-5 4-7 7-8 8-9 8-11 9-10

ring bonds :

1-2 1-4 2-3 3-4 exact/norm bonds :

1-2 1-4 1-6 2-3 3-4 3-5 4-7 7-8 8-9 8-11

exact bonds :

isolated ring systems :

containing 1 :

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS

10:CLASS 11:CLASS

L1 STRUCTURE UPLOADED

=> d l1

L1 HAS NO ANSWERS

L1

Structure attributes must be viewed using STN Express query preparation.

=> s 11

SAMPLE SEARCH INITIATED 10:59:11 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 0 TO ITERATE

100.0% PROCESSED

0 ITERATIONS

0 ANSWERS

05/18/2007

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

0 ANSWERS

PROJECTED ITERATIONS: 0 TO 0

PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

=> s l1 sss full

FULL SEARCH INITIATED 10:59:18 FILE 'REGISTRY'

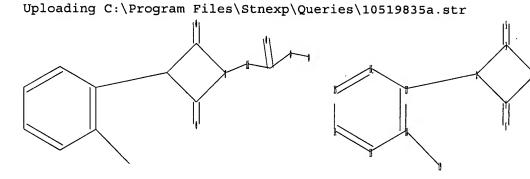
FULL SCREEN SEARCH COMPLETED - 1 TO ITERATE

100.0% PROCESSED 1 ITERATIONS

SEARCH TIME: 00.00.01

L3 0 SEA SSS FUL L1

=>



chain nodes :

5 6 7 8 9 10 11 19

ring nodes :

1 2 3 4 13 14 15 16 17 18

chain bonds :

1-6 2-17 3-5 4-7 7-8 8-9 8-11 9-10 18-19

ring bonds :

1-2 1-4 2-3 3-4 13-14 13-18 14-15 15-16 16-17 17-18

exact/norm bonds :

1-2 1-4 1-6 2-3 3-4 3-5 4-7 7-8 8-9 8-11

exact bonds :

2-17 9-10 18-19

normalized bonds :

13-14 13-18 14-15 15-16 16-17 17-18

isolated ring systems :

containing 1 :

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS 10:CLASS 11:CLASS 13:Atom 14:Atom 15:Atom 16:Atom 17:Atom 18:Atom 19:CLASS

L4 STRUCTURE UPLOADED

=> d 14

05/18/2007

L4 HAS NO ANSWERS

L4 STR

Structure attributes must be viewed using STN Express query preparation.

=> s 14

SAMPLE SEARCH INITIATED 11:02:05 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 1 TO ITERATE

100.0% PROCESSED 1 ITERATIONS

0 ANSWERS

ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 1 TO 80

PROJECTED ANSWERS: 0 TO 0

L5 0 SEA SSS SAM L4

=> s l4 sss full

L6

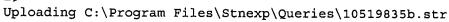
FULL SEARCH INITIATED 11:02:11 FILE 'REGISTRY'

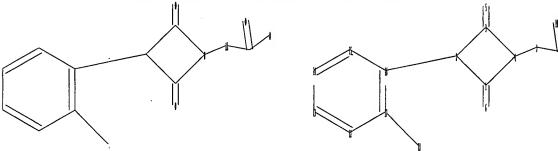
FULL SCREEN SEARCH COMPLETED - 5 TO ITERATE

100.0% PROCESSED 5 ITERATIONS

SEARCH TIME: 00.00.01

0 SEA SSS FUL L4





chain nodes :

5 6 7 8 9 10 18

ring nodes :

1 2 3 4 12 13 14 15 16 17

chain bonds :

1-6 2-16 3-5 4-7 7-8 8-9 8-10 17-18

ring bonds :

1-2 1-4 2-3 3-4 12-13 12-17 13-14 14-15 15-16 16-17

exact/norm bonds :

1-2 1-4 1-6 2-3 3-4 3-5 4-7 7-8 8-9 8-10

exact bonds : 2-16 17-18

normalized bonds :

12-13 12-17 13-14 14-15 15-16 16-17

isolated ring systems :

containing 1 :

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS 10:CLASS 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:Atom 18:CLASS

L7 STRUCTURE UPLOADED

=> d 17

L7 HAS NO ANSWERS

L7 STR

Structure attributes must be viewed using STN Express query preparation.

=> s 17

SAMPLE SEARCH INITIATED 11:03:26 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 29 TO ITERATE

100.0% PROCESSED SEARCH TIME: 00.00.01 29 ITERATIONS

0 ANSWERS

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 257 TO 903

PROJECTED ANSWERS: 0 TO

L8 0 SEA SSS SAM L7

=> s 17 sss full FULL SEARCH INITIATED 11:03:34 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED - 637 TO ITERATE

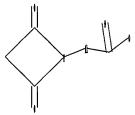
100.0% PROCESSED 637 ITERATIONS

SEARCH TIME: 00.00.01

0 ANSWERS

L9 0 SEA SSS FUL L7

=>
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chain nodes : 5 6 7 8 9 10

ring nodes : 1 2 3 4 chain bonds :

1-6 3-5 4-7 7-8 8-9 8-10

ring bonds : 1-2 1-4 2-3 3-4 exact/norm bonds :

1-2 1-4 1-6 2-3 3-4 3-5 4-7 7-8 8-9 8-10

isolated ring systems :

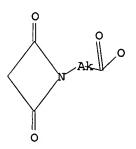
containing 1 :

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS 10:CLASS

L10 STRUCTURE UPLOADED

=> d 110 L10 HAS NO ANSWERS L10 STR



Structure attributes must be viewed using STN Express query preparation.

=> s 110

SAMPLE SEARCH INITIATED 11:05:17 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 351 TO ITERATE

100.0% PROCESSED

351 ITERATIONS

0 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 5896 TO 8144

PROJECTED ANSWERS: 0 TO

L11

0 SEA SSS SAM L10

=> s 110 sss full FULL SEARCH INITIATED 11:05:24 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED - 6664 TO ITERATE

100.0% PROCESSED 6664 ITERATIONS SEARCH TIME: 00.00.01

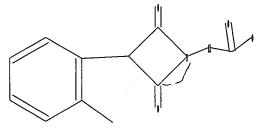
11 ANSWERS

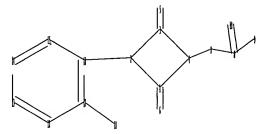
L12

11 SEA SSS FUL L10

=>

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chain nodes :

5 6 7 8 9 10 18

ring nodes :

1 2 3 4 12 13 14 15 16 17

chain bonds :

1-6 2-16 3-5 4-7 7-8 8-9 8-10 17-18

ring bonds :

1-2 1-4 2-3 3-4 12-13 12-17 13-14 14-15 15-16 16-17

exact/norm bonds :

1-2 1-4 1-6 2-3 3-4 3-5 4-7 7-8 8-9 8-10

exact bonds : 2-16 17-18

normalized bonds :

12-13 12-17 13-14 14-15 15-16 16-17

isolated ring systems :

containing 1 : 12 :

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS 10:CLASS 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:Atom 18:CLASS

L13 STRUCTURE UPLOADED

=> d 113

L13 HAS NO ANSWERS L13

Structure attributes must be viewed using STN Express query preparation.

ANSWERS -

=> s 113

SAMPLE SEARCH INITIATED 11:07:11 FILE 'REGISTRY' SAMPLE SCREEN SEARCH COMPLETED -4 TO ITERATE

100.0% PROCESSED

4 ITERATIONS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

> BATCH **COMPLETE**

PROJECTED ITERATIONS:

4 TO 200 PROJECTED ANSWERS: 0 TO

L14 0 SEA SSS SAM L13

=> s l13 sss full

FULL SEARCH INITIATED 11:07:19 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED -83 TO ITERATE

100.0% PROCESSED 83 ITERATIONS

SEARCH TIME: 00.00.01

L15 0 SEA SSS FUL L13

(FILE 'HOME' ENTERED AT 10:58:33 ON 18 MAY 2007)

FILE 'REGISTRY' ENTERED AT 10:58:47 ON 18 MAY 2007

L1 STRUCTURE UPLOADED

L2 0 S L1

L3 0 S L1 SSS FULL

L4 STRUCTURE UPLOADED

L5 0 S L4

05/18/2007

=> d his

Page 9

L6	0 S L4 SSS FULL
L7	STRUCTURE UPLOADED
L8	0 S L7
L9	0 S L7 SSS FULL
L10	STRUCTURE UPLOADED
L11	0 S L10
L12	11 S L10 SSS FULL
L13	STRUCTURE UPLOADED
L14	0 S L13
L15	0 S L13 SSS FULL

=> FIL HCAPLUS

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 865.00 865.21

FULL ESTIMATED COST

FILE 'HCAPLUS' ENTERED AT 11:07:56 ON 18 MAY 2007 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2007 AMERICAN CHEMICAL SOCIETY (ACS)

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FILE COVERS 1907 - 18 May 2007 VOL 146 ISS 22 FILE LAST UPDATED: 15 May 2007 (20070515/ED)

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This file contains CAS Registry Numbers for easy and accurate

=> s l12 L16 7 L12

=> d l16 ibib abs hitstr tot

L16 ANSWER 1 OF 7 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1992:469674 HCAPLUS

DOCUMENT NUMBER: 117:69674

TITLE: A facile synthesis of azetidine-2,4-diones

AUTHOR(S): Bari, S. S.; Trehan, I. R.; Sharma, A. K.; Manhas, M.

s.

CORPORATE SOURCE: Dep. Chem., Panjab Univ., Chandigarh, India

SOURCE: Synthesis (1992), (5), 439-42

CODEN: SYNTBF; ISSN: 0039-7881

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 117:69674

GI

AB Azetidine-2,4-diones, e.g. I, can be conveniently synthesized by mild oxidative hydrolysis of 4,4-bis(alkylthio)azetidin-2-ones, e.g. II, using N-bromosuccinimide.

IT 142389-08-0P 142389-09-1P

RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of)

RN 142389-08-0 HCAPLUS

CN 1-Azetidineacetic acid, 2,4-dioxo-3-phenoxy-, ethyl ester (9CI) (CA INDEX NAME)

RN 142389-09-1 HCAPLUS

CN 1-Azetidineacetic acid, 3-(1,3-dihydro-1,3-dioxo-2H-isoindol-2-yl)-2,4-dioxo-, ethyl ester (9CI) (CA INDEX NAME)

L16 ANSWER 2 OF 7 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1989:407106 HCAPLUS

DOCUMENT NUMBER:

111:7106

TITLE:

Studies related to penicillins. Part 26. Conversion of potassium benzylpenicillinate into 1-substituted derivatives of 3-phenylacetamidoazetidine-2,4-dione

AUTHOR(S):

CORPORATE SOURCE:

Kaura, Arun C.; Stoodley, Richard J.
Dep. Org. Chem., Univ. Newcastle upon Tyne, Newcastle

upon Tyne, NE1 7RU, UK

SOURCE:

Journal of the Chemical Society, Perkin Transactions
1: Organic and Bio-Organic Chemistry (1972-1999)

(1988), (10), 2813-20

CODEN: JCPRB4; ISSN: 0300-922X

DOCUMENT TYPE:

Journal

LANGUAGE:

English

OTHER SOURCE(S):

CASREACT 111:7106

A new method for the synthesis of azetidine-2,4-diones, involving a · AB Norrish Type II photoreaction of 4-pyruvoylazetidin-2-ones, has been devised. The process features in two strategies in which potassium benzylpenicillinate is converted into azetidine-2,4-diones I (R = Me, CH2C6H4NO2-4, CHCOMe). In one strategy, oxadiazabicyclo[3.2.0]heptenones II were treated with pyruvic acid to give the (3R,4R)-3-phenylacetamido-4pyruvoyloxyazetidin-2-ones (III), photolysis converted III into I. II were derived from (2R,3S)-4-oxo-3-phenylacetamidoazetidine-2-sulfinic acids by a novel oxidative desulfinylation induced by Pb(OAc)4. In the second strategy, (1S,5R)-analogs of II were treated with pyruvic acid to give (3S,4S)-3-phenylacetamido-4-pyruvoyloxyazetidin-2-ones. Photolysis with loss of MeSH gave I (R = Me). I (R = CH2OMe) was transformed into I (R = H) by CF3CO2H. I (R = H) lacked antibacterial activity and β -lactamase-inhibitory properties.

IT 121003-02-9P

> RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation and ester hydrolysis of)

RN 121003-02-9 HCAPLUS

CN 1-Azetidineacetic acid, α -(1-methylethylidene)-2,4-dioxo-3-[(phenylacetyl)amino]-, methoxymethyl ester (9CI) (CA INDEX NAME)

IT 71840-43-2P 121002-97-9P

RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of)

RN71840-43-2 HCAPLUS

CN 1-Azetidineacetic acid, α -(1-methylethylidene)-2,4-dioxo-3-[(phenylacetyl)amino]-, methyl ester (9CI) (CA INDEX NAME)

$$\begin{array}{c|c} & \text{Me}_2\text{C} & \text{O} \\ & \parallel & \parallel \\ & \text{C-C-OMe} \\ \\ \text{Ph-CH}_2\text{-C-NH} & \text{O} \\ \end{array}$$

RN 121002-97-9 HCAPLUS

CN 1-Azetidineacetic acid, α -(1-methylethylidene)-2,4-dioxo-3-[(phenylacetyl)amino]-, (4-nitrophenyl)methyl ester (9CI) (CA INDEX NAME)

IT 121003-03-0P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation, esterification, and bactericidal activity of)

RN 121003-03-0 HCAPLUS

CN 1-Azetidineacetic acid, α -(1-methylethylidene)-2,4-dioxo-3-[(phenylacetyl)amino]- (9CI) (CA INDEX NAME)

L16 ANSWER 3 OF 7 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1982:6476 HCAPLUS

DOCUMENT NUMBER: 96:6476

TITLE: 5,6-Dehydropenicillins

INVENTOR(S): Re, Luciano; Brant, Alberto; Bassignani, Luciano

PATENT ASSIGNEE(S): Snamprogetti SpA, Italy

SOURCE: U.S., 9 pp. Division of U.S. Ser. No. 949,546,

abandoned.

CODEN: USXXAM

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

	_				
US 4288366	Α	19810908	US 1979-58945		19790719
US 4133807	Α	19790109	US 1977-769527		19770217
GB 1572140	A	19800723	GB 1979-1747		19770222
US 4353825	A	19821012	US 1981-236029		19810219
DK 8101952	Α.	19810501	DK 1981-1952		19810501
DK 8105587	A	19811216	DK 1981-5587		19811216
PRIORITY APPLN. INFO.:			US 1978-949546	A3	19781010
			US 1977-769527	A3	19770217
			IT 1976-20451	Α	19760223
			IT 1976-20457		19760223
	•		DK 1977-771	Α	19770222
			GB 1977-7500	A	19770222
			US 1979-58945	A3	19790719

OTHER SOURCE(S):

MARPAT 96:6476

GI

AB Irradiation of I gave II which was cyclized to pencillanate III.

ΙT 79977-00-7P

> RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of)

RN 79977-00-7 HCAPLUS

CN 1-Azetidineacetic acid, 3-(acetylamino)- α -(1-methylethylidene)-2,4dioxo-, methyl ester (9CI) (CA INDEX NAME)

L16 ANSWER 4 OF 7 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1980:180878 HCAPLUS

DOCUMENT NUMBER:

92:180878

TITLE: Properties and reactions of 4-thioxo-2-azetidinones AUTHOR(S): Bachi, Mario D.; Goldberg, Ora; Gross, Akiva; Vaya,

Jacob

CORPORATE SOURCE: Dep. Org. Chem., Weizmann Inst. Sci., Rehovot, Israel

SOURCE: Journal of Organic Chemistry (1980), 45(8), 1481-5

CODEN: JOCEAH; ISSN: 0022-3263

DOCUMENT TYPE:

LANGUAGE:

Journal English

GΙ

AB 4-Thioxo-2-azetidinones I-IV appear to be suitable substrates for contrasting the chemical of the C:O and C:S linkages. Hydrolysis and alcoholysis occur selectively at the carbonyl bond while 1,3-dipolar reagents like diazoalkanes and ozone, as well as carbenes, attack exclusively at the thiocarbonyl function. The 4-alkylidene-2-azetidinones, e.g. I, were obtained from the 4-thioxo-2-azetidinones III or IV and 2-diazopropane, diphenyldiazomethane, or Et diazomalonate. The reactions with 2-diazopropane involved the formation of thiadiazolines from which the S and N elements were extruded. The reactions with the last two reagents which were performed in the presence of Rh(OAc)2 involved carbene intermediates.

IT 69939-41-9P

RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of)

RN 69939-41-9 HCAPLUS

CN 1-Azetidineacetic acid, 3-(1,3-dihydro-1,3-dioxo-2H-isoindol-2-yl)- α -(1-methylethylidene)-2,4-dioxo-, methyl ester (9CI) (CA INDEX NAME)

L16 ANSWER 5 OF 7 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1980:76192 HCAPLUS

DOCUMENT NUMBER: 92:76192

TITLE: Preparation of 3-phenylacetamidoazetidine-2,4-diones

AUTHOR (S): Kaura, Arun C.; Stoodley, Richard J.

CORPORATE SOURCE: Dep. Org. Chem., Univ. Newcastle upon Tyne, Newcastle

upon Tyne, UK

SOURCE: Journal of the Chemical Society, Chemical

Communications (1979), (7), 344-5

CODEN: JCCCAT; ISSN: 0022-4936

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 92:76192

GI

- Irradiation of pyruvates I [R = C(CO2Me) : CMe2, Q, R1 = H, R2 = O2CCOMe], AB prepared (72% in the former case) by treating oxazoloazetidinones II (R as before) with pyruvic acid, in benzene gave diones I (R as before, R1R2 = O) in 87 and 43% yields, resp.
- IT 71840-43-2P

RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of)

RN71840-43-2 HCAPLUS

CN1-Azetidineacetic acid, α -(1-methylethylidene)-2,4-dioxo-3-[(phenylacetyl)amino]-, methyl ester (9CI) (CA INDEX NAME)

$$\begin{array}{c|c} & \text{Me}_2C & \text{O} \\ & \parallel & \parallel \\ & \text{C-C-OMe} \\ \\ \text{Ph-CH}_2\text{-C-NH} & \text{O} \\ \end{array}$$

L16 ANSWER 6 OF 7 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1979:168383 HCAPLUS

DOCUMENT NUMBER: 90:168383

TITLE: Reactions of 4-thioxo-2-azetidinones: synthesis of a

2,4-azetidinedione and of 4-alkylidene-2-azetidinones AUTHOR(S): Bachi, Mario D.; Goldberg, Ora; Gross, Akiva

CORPORATE SOURCE: Dep. Org. Chem., Weizmann Inst. Sci., Rehovot, Israel

SOURCE: Tetrahedron Letters (1978), (43), 4167-70

CODEN: TELEAY; ISSN: 0040-4039

DOCUMENT TYPE: Journal

LANGUAGE: Coulinat

GI

$$\begin{array}{c|c}
 & Z \\
 & NC (CO_2Me) = CMe_2
\end{array}$$

No
$$(CO_2Me) = CMe_2$$

AB Thioxoazetidinone I (Z = S), prepared from sulfoxide II by sequential ring cleavage with CH2:CHCO2Me, isomerization, and thermal elimination reaction, on ozonolysis gave 85% I (Z = O). I (Z = O) is the 1st example of a malonimide bearing an imido group at C-3. Reaction of I (Z = S) with

MeCHN2 gave 45% thiirane III (X = bond, R = H), and with EtCHN2 gave 75% III (X = N:N, R = Me) (IV). III (X = bond, R = Me) was obtained (quant.) on decomposition of IV at ambient temperature for 5 days. I (Z = CHMe, CMe2)

were

obtained (95%) by desulfuration of III (X = bond, R = H, Me, resp.) with Ph3P in C6H6.

69939-41-9P IT

RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of)

69939-41-9 HCAPLUS RN

CN 1-Azetidineacetic acid, 3-(1,3-dihydro-1,3-dioxo-2H-isoindol-2-yl)- α -(1-methylethylidene)-2,4-dioxo-, methyl ester (9CI) (CA INDEX NAME)

L16 ANSWER 7 OF 7 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1972:112987 HCAPLUS

DOCUMENT NUMBER: 76:112987

TITLE: Synthesis of malonimide derivatives as potential

penicillin analogs

AUTHOR(S): Golik, Uri

CORPORATE SOURCE: Dep. Chem., Weizmann Inst. Sci., Rehovot, Israel

SOURCE: Journal of Heterocyclic Chemistry (1972), 9(1), 21-4

CODEN: JHTCAD; ISSN: 0022-152X

DOCUMENT TYPE: Journal LANGUAGE: English

GI For diagram(s), see printed CA Issue.

Malonimidoacetic acid derivs. (I, II) were synthesized as potential AB penicillin analogs, but they failed to inhibit the growth of bacteria when

tested in vitro against a range of gram-pos. and gram-neg. microorganisms. 35359-51-4P 35359-54-7P 35359-55-8P

IT

RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of)

RN 35359-51-4 HCAPLUS

CN 1-Azetidineacetic acid, 3,3-diethyl-2,4-dioxo-, ethyl ester (9CI) INDEX NAME)

RN 35359-54-7 HCAPLUS CN 1-Azetidineacetic acid, 3,3-diethyl-2,4-dioxo-, phenylmethyl ester (9CI) (CA INDEX NAME)

$$\begin{array}{c} O \\ \parallel \\ CH_2-C-O-CH_2-Ph \\ \\ Et \end{array}$$

RN 35359-55-8 HCAPLUS

1-Azetidineacetic acid, 3,3-diethyl-2,4-dioxo- (9CI) (CA INDEX NAME) CN

$$\begin{array}{c|c} \mathsf{CH}_2-\mathsf{CO}_2\mathsf{H} \\ \\ \mathsf{Et} & \mathsf{O} \end{array}$$

=> s imidoalkanpercarboxylic acids

0 IMIDOALKANPERCARBOXYLIC

1574229 ACIDS

L17

0 IMIDOALKANPERCARBOXYLIC ACIDS (IMIDOALKANPERCARBOXYLIC(W) ACIDS)

=> s imidoalkanpercarboxylic

0 IMIDOALKANPERCARBOXYLIC

L18

0 IMIDOALKANPERCARBOXYLIC

=> s imidopercarboxylic

5 IMIDOPERCARBOXYLIC

=> d l19 ibib abs hitstr tot

L19 ANSWER 1 OF 5 HCAPLUS COPYRIGHT 2007 ACS OF STN

ACCESSION NUMBER:

2005:1042379 HCAPLUS

DOCUMENT NUMBER:

143:328182

TITLE:

Dilution process for imidoperearboxylic

acids solution

INVENTOR(S): PATENT ASSIGNEE(S) Bianchi, Ugo Piero; Garaffa, Rober

SOURCE:

Solvay Solexis S.p.A. PCT Int. Appl., 22 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

. English

FAMILY ACC. NUM. COUNT:

1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE '
WO 2005090544	A1	20050929	WO 2005-EP51172	20050315

05/18/2007

Page 19

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     EP 1727887
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                           A1
                                                                      20050315
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     CN 1930278
                                 20070314
                                              CN 2005-80008194
                                                                      20050315
PRIORITY APPLN. INFO.:
                                              IT 2004-MI497
                                                                      20040316
                                              WO 2005-EP51172
                                                                      20050315
OTHER SOURCE(S):
                          MARPAT 143:328182
     A process for obtaining dilute aqueous solns. containing an amount of < 7% by
weight,
     expressed as a percentage by weight, of imidoalkanepercarboxylic acids,
     starting with concentrated aqueous compns. of the said peracids in the \beta form,
     the said concentrated compns. being obtained from the imidoalkanepercarboxylic
     acids in the \alpha-crystal form, comprises the following steps: (I)
     dilution of the concentrated aqueous composition of imidoalkanepercarboxylic
acids (C) with
     an aqueous solution (D) having a pH of between 2 and 5, in a (C):(D)
proportion,
     expressed in parts by weight, of between 0.1:10 and 10:0.2, working at temps.
     of between 4° and 30°; (II) application to the dilute aqueous
     composition obtained in (I) of a shear force off at least 5000 s-1, until a
     constant dynamic viscosity is obtained; (III) dilution of the dilute aqueous
composition
     treated in (II) to obtain an imidoalkanepercarboxylic acid concentration of <
     by weight; (IV) optionally, final homogenization of the composition Such
dilute aqueous
     compns. are used in detergency applications, in disinfection applications,
     and are applied by means of dispersers (triggers).
REFERENCE COUNT:
                                THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS
                          5
                                RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L19 ANSWER 2 OF 5 HCAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER:
                          2004:1127241 HCAPLUS
DOCUMENT NUMBER:
                          142:76602
TITLE:
                          Manufacture of storage-stable percarboxylic acid-based
                          polyelectrolyte capsule system
INVENTOR(S):
                          Schmiedel, Peter; Buzzacchi, Matteo; Kaiser, Heribert;
                          Von Rybinski, Wolfgang; Orlich, Bernhard
PATENT ASSIGNEE(S):
                          Henkel Kommanditgesellschaft auf Aktien, Germany
SOURCE:
                          PCT Int. Appl:, 67 pp.
                          CODEN: PIXXD2
DOCUMENT TYPE:
                          Patent
LANGUAGE:
                          German
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
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                                 DATE
                                             APPLICATION NO.
                                                                     DATE
     WO 2004110613
                           A1
                                 20041223
                                             WO 2004-EP6169
                                                                     20040608
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Page 20

05/18/2007

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     DE 10361170
                          A1.
                                 20050105
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                                 20060803
     US 2006172909
                          A1
                                             US 2005-303060
                                                                     20051213
PRIORITY APPLN. INFO.:
                                             DE 2003-10327127
                                                                  A 20030613
                                             DE 2003-10361170
                                                                  A 20031222
                                             WO 2004-EP6169
                                                                  W 20040608
     A multilayer capsule system containing organic percarboxylic acids, especially
AB
     imidopercarboxylic acids (e.g. 6-phthalimidoperoxycaproic acid),
     useful in liquid detergents and cleaners, dental care products, hair dyes,
     and decolorants or bleaching agents, is manufactured by depositing on
     percarboxylic acid particles ≥2 different shell layers each based
     on a polyelectrolyte and/or ionic surfactant. Thus, spraying 300 g Eureco
     W (72% 6-phthalimidoperoxycaproic acid) at 50° in a fluidized bed
     with 150 mL Luviquat Care (polycation) containing 1% Sequion 10H60, drying,
     spraying with 100 mL Sokalan CP 45 containing 1% Sequion 10H60 and repeating
     the procedure gave encapsulated particles (particle size 200-2000 \mu m)
     containing 61% active substance (6-phthalimidoperoxycaproic acid).
REFERENCE COUNT:
                         . 4
                                THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS
                                RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L19 ANSWER 3 OF 5 HCAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER:
                          2004:1127240 HCAPLUS
DOCUMENT NUMBER:
                          142:76601
TITLE:
                         Manufacture of storage-stable percarboxylic acid-based
                         capsules
INVENTOR(S):
                          Schmiedel, Peter; Kaiser, Heribert; Scholl, Elke; Von
                         Rybinski, Wolfgang
PATENT ASSIGNEE(S):
                         Henkel Kommanditgesellschaft auf Aktien, Germany
SOURCE:
                         PCT Int. Appl., 55 pp.
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
                         German
LANGUAGE:
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
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                                 DATE
                                             APPLICATION NO.
                                                                     DATE
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                                 20041223
     WO 2004110612
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                                             WO 2004-EP6168
                                                                     20040608
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             NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ,
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PRIORITY APPLN. INFO.:
                                                       DE 2003-10327127
                                                                                 A 20030613
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                                                                                 A 20031222
                                                       WO 2004-EP6168
                                                                                 W 20040608
      A capsule system containing organic percarboxylic acids, especially
AB
      imidopercarboxylic acids (e.g. 6-phthalimidoperoxycaproic acid),
      useful in liquid detergents and cleaners, dental care products, hair dyes,
      and decolorants or bleaching agents, is manufactured by depositing on
      percarboxylic acid particles ≥1 inorg. acid salt. Thus, spraying
      800 g Eureco W (72% 6-phthalimidoperoxycaproic acid) with 800 g of 20% aqueous
      Na2SO4 solution containing 1% Sequion 10H60 and separating particles <2.0 mm by
      sieving gave encapsulated bleach particles containing 70.5% active substance.
      The encapsulated particles were incorporated in a liquid detergent
      formulation which was stored at 40° to show 99% and 95% bleach
      activity after 1 and 2 wk, resp., vs. 75% and 65% for similar composition
      containing unencapsulated Eureco W.
REFERENCE COUNT:
                               6
                                       THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS
                                       RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L19 ANSWER 4 OF 5 HCAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER:
                               2004:1127239 HCAPLUS
DOCUMENT NUMBER:
                               142:76600
TITLE:
                               Manufacture of gel capsules containing percarboxylic
                               acid bleach
INVENTOR(S):
                               Schmiedel, Peter; Barreleiro, Paula; Von Rybinski,
                               Wolfgang; Orlich, Bernhard
PATENT ASSIGNEE(S):
                               Henkel Kommanditgesellschaft auf Aktien, Germany
SOURCE:
                              PCT Int. Appl., 55 pp.
                               CODEN: PIXXD2
DOCUMENT TYPE:
                               Patent
LANGUAGE:
                               German
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
      PATENT NO.
                                                       APPLICATION NO.
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                                        DATE
                                                                                  DATE
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                                       2004122/3
      WO 2004110611
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                                                                                    20040608
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SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE,

AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,

DE 2003-10361084

EP 2004-739696

JP 2006-515852

US 2005-299795

20031222

20040608

20040608

20051212

20050105

20060315

IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK

20061130

20060810

A1

A1

A1

T

SN, TD, TG

DE 10361084

JP 2006527292

US 2006178285

EP 1633470

naphthalenedicarboximido; each R = H, (un)substituted alkyl, OH, CO2H,
CO2OH, CO2R1; R1 = C1-5 (un)substituted alkyl; n = 1-5], prepared by the
reaction of H2O2 with the acid or anhydride in concentrated H2SO4 or MeSO3H or
in an alkaline medium, are useful as bleaching agents, especially in the
laundering

of fabrics at low temps. Phthalimidoacetic acid in MeSO3H was treated with H2O2 to give phthalimidoperacetic acid, which was used with a laundry detergent at 40° for the bleaching of cotton fabrics stained with red wine.

=> FIL REGISTRY

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STRUCTURE FILE UPDATES: 17 MAY 2007 HIGHEST RN 935249-87-9 DICTIONARY FILE UPDATES: 17 MAY 2007 HIGHEST RN 935249-87-9

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH December 2, 2006

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http://www.cas.org/support/stngen/stndoc/properties.html

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ring nodes :

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chain bonds :

1-6 2-12 3-5 13-14

ring bonds :

1-2 1-4 2-3 3-4 8-9 8-13 9-10 10-11 11-12 12-13

exact/norm bonds :

1-2 1-4 1-6 2-3 3-4 3-5

exact bonds : 2-12 13-14

normalized bonds :

8-9 8-13 9-10 10-11 11-12 12-13

isolated ring systems :

containing 1 : 8 :

Match level :

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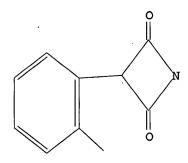
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L20 HAS NO ANSWERS

L20

STR



Structure attributes must be viewed using STN Express query preparation.

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SAMPLE SCREEN SEARCH COMPLETED - 4 TO ITERATE

100.0% PROCESSED 4 ITERATIONS

1 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 4 TO 200 PROJECTED ANSWERS: 1 TO 80

L21 1 SEA SSS SAM L20

=> s 120 sss full

05/18/2007

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100.0% PROCESSED

83 ITERATIONS

2 ANSWERS

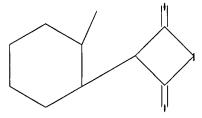
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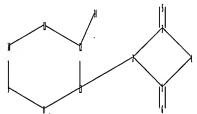
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exact bonds :

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isolated ring systems :

containing 1 : 8 :

Match level :

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12:Atom 13:Atom 14:CLASS

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PRIORITY APPLN. INFO.:

DE 2003-10327127 A 20030613 DE 2003-10361084 A 20031222 WO 2004-EP6167 W 20040608

Storage-stable gel capsules containing organic percarboxylic acids, especially AB imidopercarboxylic acids (e.g. 6-phthalimidoperoxycaproic acid), useful especially in liquid detergents and cleaners, dental care products, hair dyes, and decolorants or bleaching compns. for tech. applications, were manufactured by incorporating solid percarboxylic acid particles in a gel matrix obtained by solidification and/or gelling of an oil phase m. <35°, e.g., a paraffin oil, glyceride, vegetable oil, etc., and containing a gel-forming agent and stabilizing agent, e.g., a block copolymer. For example, adding 2% Kraton G 1650 to low-viscosity paraffin oil at 70° with stirring gave a homogeneous solution which was cooled to 40°, 60% (based on paraffin oil + Kraton) of homogenized Eureco W (72% 6-phthalimidoperoxycaproic acid) was added with stirring, the mixture was poured in 5 vols. of H2O with stirring, and cooled. The encapsulated 6-phthalimidoperoxycaproic acid particles which settled at the bottom were separated and the fraction <1 mm incorporated (6%) in a liquid detergent composition

which was stored at 40° to give 100.0% and 99.2% active O content after 1 and 2 wk, resp., vs. 87.5 and 80.0% for similar detergent containing non-capsulated Eureco W.

REFERENCE COUNT:

THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L19 ANSWER 5 OF 5 HCAPLUS COPYRIGHT 2007 ACS on STN

4

ACCESSION NUMBER:

1990:80000 HCAPLUS

DOCUMENT NUMBER: TITLE:

112:80000
Bleaching agents comprising aromatic imide

percarboxylic acids and their preparation and use

INVENTOR(S):

Venturello, Carlo; Cavallotti, Caludio; Burzio, Fulvio

PATENT ASSIGNEE(S):

Ausimont S.p.A., Italy

SOURCE:

Eur. Pat. Appl., 11 pp. CODEN: EPXXDW

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.		DATE
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EP 325289	A1	19890726	EP 1989-101003		19890120
EP 325289	B1	19930901			
R: AT, BE, CH,	DE, ES	, FR, GB,	LI, NL, SE		
AU 8928678	A	19890720	AU 1989-28678		19890120
AU 614322	B2	19910829			
JP 02001473	Α	19900105	JP 1989-11742		19890120
JP 2786222	B2	19980813			
AT 93908	\cdot T	19930915	AT 1989-101003		19890120
ES 2058348	Т3	19941101	ES 1989-101003		19890120
CA 1340680	С	19990727	CA 1989-588845		19890120
BR 8900265	A	19890919	BR 1989-265		19890123
US_5520844	A	19960528	US 1991-637479		19910104
US_568,8434	A	19971118	US 1996-593655		19960129
PRIORETY APPLN. INFO.:			IT 1988-19131	Α	19880120
			US 1989-298918	B1	19890119
			EP 1989-101003	Α	19890120
			US 1991-637479	A3	19910104

OTHER SOURCE(S): MARPAT 112:80000

AB Peroxy acids X(CHR)nCO2OH [X = (un)substituted phthalimido or

Connecting via Winsock to STN

Welcome to STN International! Enter x:x

LOGINID: SSSPTA1626GMS

PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

```
NEWS 1
                 Web Page for STN Seminar Schedule - N. America
                 CHEMLIST enhanced with New Zealand Inventory of Chemicals
NEWS 2 JAN 08
                 CA/CAplus Company Name Thesaurus enhanced and reloaded
NEWS 3 JAN 16
                 IPC version 2007.01 thesaurus available on STN
NEWS 4 JAN 16
                 WPIDS/WPINDEX/WPIX enhanced with IPC 8 reclassification data
NEWS 5 JAN 16
NEWS 6 JAN 22
                 CA/CAplus updated with revised CAS roles
NEWS 7 JAN 22
                 CA/CAplus enhanced with patent applications from India
                 PHAR reloaded with new search and display fields
NEWS 8 JAN 29
NEWS 9 JAN 29
                 CAS Registry Number crossover limit increased to 300,000 in
                 multiple databases
                PATDPASPC enhanced with Drug Approval numbers
NEWS 10 FEB 15
NEWS 11 FEB 15
                RUSSIAPAT enhanced with pre-1994 records
                KOREAPAT enhanced with IPC 8 features and functionality
NEWS 12 FEB 23
NEWS 13 FEB 26 MEDLINE reloaded with enhancements
NEWS 14 FEB 26 EMBASE enhanced with Clinical Trial Number field
NEWS 15 FEB 26
                TOXCENTER enhanced with reloaded MEDLINE
NEWS 16 FEB 26
                IFICDB/IFIPAT/IFIUDB reloaded with enhancements
NEWS 17 FEB 26 CAS Registry Number crossover limit increased from 10,000
                 to 300,000 in multiple databases
NEWS 18 MAR 15
                WPIDS/WPIX enhanced with new FRAGHITSTR display format
NEWS 19 MAR 16
                CASREACT coverage extended
NEWS 20 MAR 20
                MARPAT now updated daily
NEWS 21 MAR 22
                LWPI reloaded
NEWS 22 MAR 30
                RDISCLOSURE reloaded with enhancements
NEWS 23 APR 02
                JICST-EPLUS removed from database clusters and STN
NEWS 24 APR 30
                GENBANK reloaded and enhanced with Genome Project ID field
NEWS 25 APR 30
                CHEMCATS enhanced with 1.2 million new records
       APR 30
NEWS 26
                CA/CAplus enhanced with 1870-1889 U.S. patent records
NEWS 27
                INPADOC replaced by INPADOCDB on STN
        APR 30
NEWS 28
        MAY 01
                New CAS web site launched
NEWS 29
                CA/CAplus Indian patent publication number format defined
        MAY 08
NEWS 30
        MAY 14
                RDISCLOSURE on STN Easy enhanced with new search and display
                 fields
             NOVEMBER 10 CURRENT WINDOWS VERSION IS V8.01c, CURRENT
NEWS EXPRESS
             MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
             AND CURRENT DISCOVER FILE IS DATED 25 SEPTEMBER 2006.
NEWS HOURS
              STN Operating Hours Plus Help Desk Availability
NEWS LOGIN
              Welcome Banner and News Items
NEWS IPC8
             For general information regarding STN implementation of IPC 8
```

Enter NEWS followed by the item number or name to see news on that specific topic.

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FILE 'HOME' ENTERED AT 14:33:37 ON 18 MAY 2007

=>
Uploading

THIS COMMAND NOT AVAILABLE IN THE CURRENT FILE Do you want to switch to the Registry File? Choice (Y/n):

Switching to the Registry File...

Some commands only work in certain files. For example, the EXPAND command can only be used to look at the index in a file which has an index. Enter "HELP COMMANDS" at an arrow prompt (=>) for a list of commands which can be used in this file.

=> FILE REGISTRY

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 0.21 0.21

FULL ESTIMATED COST

FILE 'REGISTRY' ENTERED AT 14:33:53 ON 18 MAY 2007 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
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STRUCTURE FILE UPDATES: 17 MAY 2007 HIGHEST RN 935249-87-9 DICTIONARY FILE UPDATES: 17 MAY 2007 HIGHEST RN 935249-87-9

New CAS Information Use Policies, enter HELP USAGETERMS for details.

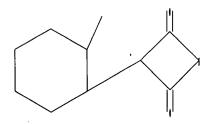
TSCA INFORMATION NOW CURRENT THROUGH December 2, 2006

Please note that search-term pricing does apply when conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/support/stngen/stndoc/properties.html

Uploading C:\Program Files\Stnexp\Queries\10519835f.str



chain nodes : 5 6 14 ring nodes :

1 2 3 4 8 9 10 11 12 13

chain bonds :

1-6 2-13 3-5 12-14

ring bonds :

1-2 1-4 2-3 3-4 8-9 8-13 9-10 10-11 11-12 12-13

exact/norm bonds :

1-2 1-4 1-6 2-3 3-4 3-5

exact bonds : .

2-13 8-9 8-13 9-10 10-11 11-12 12-13 12-14

isolated ring systems :

containing 1 : 8 :

Match level :

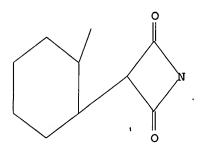
1:Atom 2:Atom 3:Atom 4:Atom 5:CLASS 6:CLASS 8:Atom 9:Atom 10:Atom 11:Atom 12:Atom 13:Atom 14:CLASS

L1 STRUCTURE UPLOADED

=> d l1

L1 HAS NO ANSWERS

L1 STR



Structure attributes must be viewed using STN Express query preparation.

=> s l1

SAMPLE SEARCH INITIATED 14:34:06 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 0 TO ITERATE

100.0% PROCESSED

0 ITERATIONS

0 ANSWERS

05/18/2007

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

L2 0 SEA SSS SAM L1

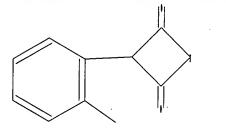
=> s l1 sss full FULL SEARCH INITIATED 14:34:12 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED - 52 TO ITERATE

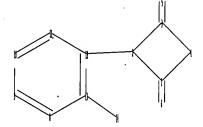
100.0% PROCESSED 52 ITERATIONS 0 ANSWERS

SEARCH TIME: 00.00.01

L3 0 SEA SSS FUL L1

Uploading C:\Program Files\Stnexp\Queries\10519835e.str





chain nodes: 5 6 14

ring nodes :

=>

1 2 3 4 8 9 10 11 12 13

chain bonds :

1-6 2-12 3-5 13-14

ring bonds :

1-2 1-4 2-3 3-4 8-9 8-13 9-10 10-11 11-12 12-13

exact/norm bonds :

1-2 1-4 1-6 2-3 3-4 3-5

exact bonds : 2-12 13-14

normalized bonds :

8-9 8-13 9-10 10-11 11-12 12-13

isolated ring systems :

containing 1 : 8 :

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:CLASS 6:CLASS 8:Atom 9:Atom 10:Atom 11:Atom 12:Atom 13:Atom 14:CLASS

L4 STRUCTURE UPLOADED

=> d 14 L4 HAS NO ANSWERS

T.4

STR

Structure attributes must be viewed using STN Express query preparation.

=> s 14

SAMPLE SEARCH INITIATED 14:36:03 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 4 TO ITERATE

100.0% PROCESSED

4 ITERATIONS

1 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS:

4 TO 200

PROJECTED ANSWERS:

1 TO 80

L5

1 SEA SSS SAM L4

=> s l4 sss full

FULL SEARCH INITIATED 14:36:09 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED -

83 TO ITERATE

100.0% PROCESSED

83 ITERATIONS

2 ANSWERS

SEARCH TIME: 00.00.01

L6

2 SEA SSS FUL L4

=> FIL HCAPLUS

COST IN U.S. DOLLARS

SINCE FILE TOTAL

ENTRY SESSION

FULL ESTIMATED COST

345.10 345.31

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FILE COVERS 1907 - 18 May 2007 VOL 146 ISS 22 FILE LAST UPDATED: 15 May 2007 (20070515/ED)

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This file contains CAS Registry Numbers for easy and accurate

=> s 16

L7 1 L6

=> d 17 ibib abs hitstr tot

L7 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1940:6252 HCAPLUS

DOCUMENT NUMBER: 34:6252

ORIGINAL REFERENCE NO.: 34:1012i,1013a-c

TITLE: Azetidine derivatives. I. 2,4-Diketo-3-hydroxy-3-

arylazetidines

AUTHOR(S): Riebsomer, J. L.; Burkett, Howard; Hodgson, Thomas;

Senour, Fred

SOURCE: Journal of the American Chemical Society (1939), 61,

3491-3

CODEN: JACSAT; ISSN: 0002-7863

DOCUMENT TYPE:

Journal Unavailable

LANGUAGE: Unavailable GI For diagram(s), see printed CA Issue.

Ph(OH)C(CO2Et)2 (25 g.) and 7.5 g. Na in 80 cc. absolute EtOH, heated to AB 115-20° and treated with 10 g. CO(NH2)2, stirred at 115-20° for 5 hrs., 10 g. CO(NH2)2 again added and the stirring and heating continued for about 11 hrs., the solution evaporated to dryness, the product dissolved in 100 cc. H2O, 50 g. ice and 10% HCl added until the solution is acid, and the solution extracted with Et2O, give 17% of 2,4-diketo-3-hydroxy-3phenylazetidine, PhC(OH).CO.NH.CO, m. 107.5-8°; NH3 may be bubbled into the mixture instead of the addition of CO(NH2)2, the yield being about 3%. Similarly prepared were the p-tolyl derivative, m. 131° (9%); the p-ethylphenyl derivative, m. 105-6° (24%); the 2,5-dimethylphenyl derivative, m. 135-6° (37%); the mesityl derivative, m. 151-2° (6%); the p-sec-butylphenyl derivative, m. 89-90° (38%). The structure of these compds. follows from their hydrolysis with NaOH to PhCH(OH)CO2H and its derivs., Ph(HO)C(CO2Na)2 being assumed as an intermediate. These compds. are inactive as hypnotics when given to rabbits and are toxic in relatively small doses.

IT 857955-59-0P, Tartronimide, α -2,5-xylyl-

857955-73-8P, Tartronimide, α -(mesityl)-

RL: PREP (Preparation) (preparation of)

RN 857955-59-0 HCAPLUS

CN Tartronimide, α -2,5-xylyl- (4CI) (CA INDEX NAME)

RN 857955-73-8 HCAPLUS CN Tartronimide, α -(mesityl)- (4CI) (CA INDEX NAME)

=> FIL REGISTRY

COST IN U.S. DOLLARS

SINCE FILE TOTAL

ENTRY SESSION

FULL ESTIMATED COST

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE TOTAL

ENTRY SESSION
CA SUBSCRIBER PRICE -0.78 -0.78

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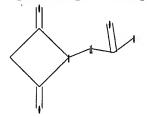
TSCA INFORMATION NOW CURRENT THROUGH December 2, 2006

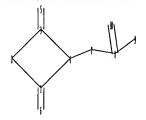
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http://www.cas.org/support/stngen/stndoc/properties.html

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chain nodes : 5 6 7 8 9 10

ring nodes : 1 2 3 4

chain bonds :

1-6 3-5 4-7 7-8 8-9 8-10

ring bonds :

1-2 1-4 2-3 3-4

exact/norm bonds :

1-2 1-4 1-6 2-3 3-4 3-5 4-7 7-8 8-9 8-10

isolated ring systems :

containing 1 :

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS

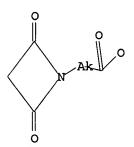
10:CLASS

L8 STRUCTURE UPLOADED

=> d 18

L8 HAS NO ANSWERS

STR



Structure attributes must be viewed using STN Express query preparation.

=> s 18

SAMPLE SEARCH INITIATED 14:39:05 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 351 TO ITERATE

100.0% PROCESSED 351 ITERATIONS

SEARCH TIME: 00.00.01

0 ANSWERS

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 5896 TO

8144 PROJECTED ANSWERS: 0 TO ٥

L9 0 SEA SSS SAM L8

=> s 18 sss full

FULL SEARCH INITIATED 14:39:12 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 6664 TO ITERATE

100.0% PROCESSED 6664 ITERATIONS 11 ANSWERS

SEARCH TIME: 00.00.01

L10 11 SEA SSS FUL L8

=> FIL HCAPLUS

COST IN U.S. DOLLARS SINCE FILE TOTAL SESSION ENTRY

FULL ESTIMATED COST 172.10 533.08

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL ENTRY SESSION

CA SUBSCRIBER PRICE 0.00 -0.78

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FILE COVERS 1907 - 18 May 2007 VOL 146 ISS 22 FILE LAST UPDATED: 15 May 2007 (20070515/ED)

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This file contains CAS Registry Numbers for easy and accurate

=> s 110

L11 7 L10

=> d l11 ibib abs hitstr tot

L11 ANSWER 1 OF 7 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1992:469674 HCAPLUS

DOCUMENT NUMBER: 117:69674

TITLE: A facile synthesis of azetidine-2,4-diones

AUTHOR (S): Bari, S. S.; Trehan, I. R.; Sharma, A. K.; Manhas, M.

CORPORATE SOURCE: Dep.

Dep. Chem., Panjab Univ., Chandigarh, India

SOURCE:

Synthesis (1992), (5), 439-42 CODEN: SYNTBF; ISSN: 0039-7881

DOCUMENT TYPE:

Journal English

LANGUAGE:
OTHER SOURCE(S):

CASREACT 117:69674

GI

AB Azetidine-2,4-diones, e.g. I, can be conveniently synthesized by mild oxidative hydrolysis of 4,4-bis(alkylthio)azetidin-2-ones, e.g. II, using N-bromosuccinimide.

IT 142389-08-0P 142389-09-1P

RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of)

RN 142389-08-0 HCAPLUS

CN 1-Azetidineacetic acid, 2,4-dioxo-3-phenoxy-, ethyl ester (9CI) (CA INDEX NAME)

RN 142389-09-1 HCAPLUS

CN 1-Azetidineacetic acid, 3-(1,3-dihydro-1,3-dioxo-2H-isoindol-2-yl)-2,4-dioxo-, ethyl ester (9CI) (CA INDEX NAME)

L11 ANSWER 2 OF 7 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1989:407106 HCAPLUS

DOCUMENT NUMBER: 111:7106

TITLE: Studies related to penicillins. Part 26. Conversion

of potassium benzylpenicillinate into 1-substituted

AUTHOR (S):

derivatives of 3-phenylacetamidoazetidine-2,4-dione

Kaura, Arun C.; Stoodley, Richard J.

CORPORATE SOURCE: Dep. Org. Chem., Univ. Newcastle upon Tyne, Newcastle

upon Tyne, NE1 7RU, UK

Journal of the Chemical Society, Perkin Transactions SOURCE:

1: Organic and Bio-Organic Chemistry (1972-1999)

(1988), (10), 2813-20 CODEN: JCPRB4; ISSN: 0300-922X

DOCUMENT TYPE: LANGUAGE:

Journal English

OTHER SOURCE(S):

CASREACT 111:7106

A new method for the synthesis of azetidine-2,4-diones, involving a AB Norrish Type II photoreaction of 4-pyruvoylazetidin-2-ones, has been devised. The process features in two strategies in which potassium benzylpenicillinate is converted into azetidine-2,4-diones I (R = Me, CH2C6H4NO2-4, CHCOMe). In one strategy, oxadiazabicyclo[3.2.0]heptenones II were treated with pyruvic acid to give the (3R,4R)-3-phenylacetamido-4pyruvoyloxyazetidin-2-ones (III), photolysis converted III into I. II were derived from (2R,3S)-4-oxo-3-phenylacetamidoazetidine-2-sulfinic acids by a novel oxidative desulfinylation induced by Pb(OAc)4. In the second strategy, (1S,5R)-analogs of II were treated with pyruvic acid to give (3S,4S)-3-phenylacetamido-4-pyruvoyloxyazetidin-2-ones. Photolysis with loss of MeSH gave I (R = Me). I (R = CH2OMe) was transformed into I (R = H) by CF3CO2H. I (R = H) lacked antibacterial activity and β-lactamase-inhibitory properties.

IT 121003-02-9P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation and ester hydrolysis of)

RN121003-02-9 HCAPLUS

CN1-Azetidineacetic acid, α -(1-methylethylidene)-2,4-dioxo-3-[(phenylacetyl)amino]-, methoxymethyl ester (9CI) (CA INDEX NAME)

TT 71840-43-2P 121002-97-9P

RL: SPN (Synthetic preparation); PREP (Preparation)

RN

(preparation of) 71840-43-2 HCAPLUS

CN 1-Azetidineacetic acid, α -(1-methylethylidene)-2,4-dioxo-3-[(phenylacetyl)amino]-, methyl ester (9CI) (CA INDEX NAME)

RN 121002-97-9 HCAPLUS

CN 1-Azetidineacetic acid, α-(1-methylethylidene)-2,4-dioxo-3[(phenylacetyl)amino]-, (4-nitrophenyl)methyl ester (9CI) (CA INDEX NAME)

IT 121003-03-0P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation, esterification, and bactericidal activity of)

RN 121003-03-0 HCAPLUS

CN 1-Azetidineacetic acid, α -(1-methylethylidene)-2,4-dioxo-3-[(phenylacetyl)amino]- (9CI) (CA INDEX NAME)

$$\begin{array}{c|c} \mathsf{CMe_2} \\ | \\ \mathsf{C-Co_2H} \\ \\ \mathsf{Ph-CH_2-C-NH} \\ \end{array}$$

L11 ANSWER 3 OF 7 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1982:6476 HCAPLUS

DOCUMENT NUMBER: 96:6476

TITLE: 5,6-Dehydropenicillins

INVENTOR(S): Re, Luciano; Brant, Alberto; Bassignani, Luciano

PATENT ASSIGNEE(S): Snamprogetti SpA, Italy

SOURCE: U.S., 9 pp. Division of U.S. Ser. No. 949,546,

abandoned.

CODEN: USXXAM

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.		DATE
				-	
US 4288366	A	19810908	US 1979-58945		19790719
US 4133807	Α	19790109	US 1977-769527		19770217
GB 1572140	Α	19800723	GB 1979-1747		19770222
US 4353825	A	19821012	US 1981-236029		19810219
DK 8101952	Α	19810501	DK 1981-1952		19810501
DK 8105587	Α	19811216	DK 1981-5587		19811216
PRIORITY APPLN. INFO.:			US 1978-949546	А3	19781010
			US 1977-769527	А3	19770217
			IT 1976-20451	Α	19760223
			IT 1976-20457		19760223
			DK 1977-771	Α	19770222
			GB 1977-7500	Α	19770222
			US 1979-58945	А3	19790719

OTHER SOURCE(S):

MARPAT 96:6476

AB Irradiation of I gave II which was cyclized to pencillanate III.

IT 79977-00-7P

RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of) RN 79977-00-7 HCAPLUS

CN 1-Azetidineacetic acid, 3-(acetylamino)- α -(1-methylethylidene)-2,4-dioxo-, methyl ester (9CI) (CA INDEX NAME)

L11 ANSWER 4 OF 7 HCAPLUS COPYRIGHT 2007 ACS on STN ACCESSION NUMBER: 1980:180878 HCAPLUS

DOCUMENT NUMBER:

92:180878

TITLE: AUTHOR(S):

Properties and reactions of 4-thioxo-2-azetidinones Bachi, Mario D.; Goldberg, Ora; Gross, Akiva; Vaya,

Jacob

CORPORATE SOURCE:

SOURCE:

Dep. Org. Chem., Weizmann Inst. Sci., Rehovot, Israel

II

Journal of Organic Chemistry (1980), 45(8), 1481-5

CODEN: JOCEAH; ISSN: 0022-3263

DOCUMENT TYPE:

LANGUAGE:

Journal English

GI

$$\begin{array}{c}
CMe_2\\
NC(CO_2Me) = CMe_2
\end{array}$$

AB 4-Thioxo-2-azetidinones I-IV appear to be suitable substrates for contrasting the chemical of the C:O and C:S linkages. Hydrolysis and alcoholysis occur selectively at the carbonyl bond while 1,3-dipolar reagents like diazoalkanes and ozone, as well as carbenes, attack exclusively at the thiocarbonyl function. The 4-alkylidene-2-azetidinones, e.g. I, were obtained from the 4-thioxo-2-azetidinones III or IV and 2-diazopropane, diphenyldiazomethane, or Et diazomalonate. The reactions with 2-diazopropane involved the formation of thiadiazolines from which the S and N elements were extruded. The reactions with the last two reagents which were performed in the presence of Rh(OAc)2 involved carbene intermediates.

IT 69939-41-9P

RN 69939-41-9 HCAPLUS

CN 1-Azetidineacetic acid, 3-(1,3-dihydro-1,3-dioxo-2H-isoindol-2-yl)- α -(1-methylethylidene)-2,4-dioxo-, methyl ester (9CI) (CA INDEX NAME)

L11 ANSWER 5 OF 7 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1980:76192 HCAPLUS

DOCUMENT NUMBER: 92:76192

TITLE: Preparation of 3-phenylacetamidoazetidine-2,4-diones

AUTHOR(S): Kaura, Arun C.; Stoodley, Richard J.

CORPORATE SOURCE: Dep. Org. Chem., Univ. Newcastle upon Tyne, Newcastle

upon Tyne, UK

SOURCE: Journal of the Chemical Society, Chemical

Communications (1979), (7), 344-5 CODEN: JCCCAT; ISSN: 0022-4936

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 92:76192

GΙ

AB Irradiation of pyruvates I [R = C(CO2Me):CMe2, Q, R1 = H, R2 = O2CCOMe], prepared (72% in the former case) by treating oxazoloazetidinones II (R as before) with pyruvic acid, in benzene gave diones I (R as before, R1R2 = O) in 87 and 43% yields, resp.

IT 71840-43-2P

RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of)

RN 71840-43-2 HCAPLUS

CN 1-Azetidineacetic acid, α -(1-methylethylidene)-2,4-dioxo-3-[(phenylacetyl)amino]-, methyl ester (9CI) (CA INDEX NAME)

L11 ANSWER 6 OF 7 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1979:168383 HCAPLUS

DOCUMENT NUMBER: 90:168383

TITLE: Reactions of 4-thioxo-2-azetidinones: synthesis of a

2,4-azetidinedione and of 4-alkylidene-2-azetidinones

AUTHOR(S): Bachi, Mario D.; Goldberg, Ora; Gross, Akiva

CORPORATE SOURCE: Dep. Org. Chem., Weizmann Inst. Sci., Rehovot, Israel

SOURCE: Tetrahedron Letters (1978), (43), 4167-70

CODEN: TELEAY; ISSN: 0040-4039

DOCUMENT TYPE: Journal

LANGUAGE: English

$$\begin{array}{c|c}
 & Z \\
 & NC (CO_2Me) = CMe_2
\end{array}$$

NOC
$$(CO_2Me) = CMe_2$$

AB Thioxoazetidinone I (Z = S), prepared from sulfoxide II by sequential ring cleavage with CH2:CHCO2Me, isomerization, and thermal elimination reaction, on ozonolysis gave 85% I (Z = O). I (Z = O) is the 1st example of a malonimide bearing an imido group at C-3. Reaction of I (Z = S) with

MeCHN2 gave 45% thiirane III (X = bond, R = H), and with EtCHN2 gave 75% III (X = N:N, R = Me) (IV). III (X = bond, R = Me) was obtained (quant.) on decomposition of IV at ambient temperature for 5 days. I (Z = CHMe, CMe2)

were

obtained (95%) by desulfuration of III (X = bond, R = H, Me, resp.) with Ph3P in C6H6.

IT 69939-41-9P

RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of)

RN 69939-41-9 HCAPLUS

CN 1-Azetidineacetic acid, 3-(1,3-dihydro-1,3-dioxo-2H-isoindol-2-yl)- α -(1-methylethylidene)-2,4-dioxo-, methyl ester (9CI) (CA INDEX NAME)

L11 ANSWER 7 OF 7 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1972:112987 HCAPLUS

DOCUMENT NUMBER: 76:112987

TITLE: Synthesis of malonimide derivatives as potential

penicillin analogs

AUTHOR(S): Golik, Uri

CORPORATE SOURCE: Dep. Chem., Weizmann Inst. Sci., Rehovot, Israel

SOURCE: Journal of Heterocyclic Chemistry (1972), 9(1), 21-4

CODEN: JHTCAD; ISSN: 0022-152X

DOCUMENT TYPE: Journal LANGUAGE: English

GI For diagram(s), see printed CA Issue.

AB Malonimidoacetic acid derivs. (I, II) were synthesized as potential penicillin analogs, but they failed to inhibit the growth of bacteria when

tested in vitro against a range of gram-pos. and gram-neg. microorganisms.

IT 35359-51-4P 35359-54-7P 35359-55-8P

RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of)

RN 35359-51-4 HCAPLUS

CN 1-Azetidineacetic acid, 3,3-diethyl-2,4-dioxo-, ethyl ester (9CI) (CA INDEX NAME)

RN 35359-54-7 HCAPLUS

CN 1-Azetidineacetic acid, 3,3-diethyl-2,4-dioxo-, phenylmethyl ester (9CI) (CA INDEX NAME)

$$\begin{array}{c} O \\ \parallel \\ CH_2 - C - O - CH_2 - Ph \\ \\ Et \end{array}$$

RN 35359-55-8 HCAPLUS

CN 1-Azetidineacetic acid, 3,3-diethyl-2,4-dioxo- (9CI) (CA INDEX NAME)

=> FIL REGISTRY
COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 52.49 585.57

FULL ESTIMATED COST

CA SUBSCRIBER PRICE

SINCE FILE TOTAL

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

ENTRY SESSION -5.46 -6.24

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STRUCTURE FILE UPDATES: 17 MAY 2007 HIGHEST RN 935249-87-9 DICTIONARY FILE UPDATES: 17 MAY 2007 HIGHEST RN 935249-87-9

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TSCA INFORMATION NOW CURRENT THROUGH December 2, 2006

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http://www.cas.org/support/stngen/stndoc/properties.html

=>

Uploading C:\Program Files\Stnexp\Queries\10519835g.str

chain nodes :

5 6 7 8 9 10 13

ring nodes:

1 2 3 4 chain bonds:

1-6 3-5 4-7 7-8 8-9 8-10 9-13

ring bonds :

1-2 1-4 2-3 3-4

exact/norm bonds :

1-2 1-4 1-6 2-3 3-4 3-5 4-7 7-8 8-9 8-10 9-13

isolated ring systems :

containing 1 :

G1:H,M

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS

10:CLASS 13:CLASS

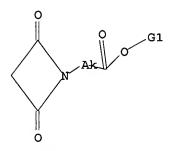
L12 STRUCTURE UPLOADED

=> d l12

L12 HAS NO ANSWERS

L12

STR



G1 H,M

Structure attributes must be viewed using STN Express query preparation.

=> s 112

SAMPLE SEARCH INITIATED 14:42:54 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 351 TO ITERATE

100.0% PROCESSED 351 ITERATIONS

0 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 5896 TO 8144

PROJECTED ANSWERS: 0 TO 0

L13 0 SEA SSS SAM L12

=> s l12 sss full

FULL SEARCH INITIATED 14:43:00 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 6664 TO ITERATE

100.0% PROCESSED 6664 ITERATIONS 2 ANSWERS

SEARCH TIME: 00.00.01

L14 2 SEA SSS FUL L12

=> FIL HCAPLUS

COST IN U.S. DOLLARS SINCE FILE TOTAL

ENTRY SESSION

FULL ESTIMATED COST 172.10 757.67

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL

CA SUBSCRIBER PRICE ENTRY SESSION 0.00 -6.24

FILE 'HCAPLUS' ENTERED AT 14:43:07 ON 18 MAY 2007 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

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FILE COVERS 1907 - 18 May 2007 VOL 146 ISS 22 FILE LAST UPDATED: 15 May 2007 (20070515/ED)

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate

=> s l14

L15 2 L14

=> d l15 ibib abs hitstr tot

L15 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

1989:407106 HCAPLUS

DOCUMENT NUMBER:

111:7106

TITLE:

Studies related to penicillins. Part 26. Conversion of potassium benzylpenicillinate into 1-substituted

derivatives of 3-phenylacetamidoazetidine-2,4-dione

AUTHOR (S):

Kaura, Arun C.; Stoodley, Richard J.

CORPORATE SOURCE:

Dep. Org. Chem., Univ. Newcastle upon Tyne, Newcastle

upon Tyne, NE1 7RU, UK

SOURCE:

Journal of the Chemical Society, Perkin Transactions

1: Organic and Bio-Organic Chemistry (1972-1999)

(1988), (10), 2813-20

CODEN: JCPRB4; ISSN: 0300-922X

DOCUMENT TYPE:

LANGUAGE:

Journal English

OTHER SOURCE(S):

CASREACT 111:7106

GI

AB A new method for the synthesis of azetidine-2,4-diones, involving a Norrish Type II photoreaction of 4-pyruvoylazetidin-2-ones, has been devised. The process features in two strategies in which potassium benzylpenicillinate is converted into azetidine-2,4-diones I (R = Me, CH2C6H4NO2-4, CHCOMe). In one strategy, oxadiazabicyclo[3.2.0]heptenones II were treated with pyruvic acid to give the (3R,4R)-3-phenylacetamido-4-pyruvoyloxyazetidin-2-ones (III), photolysis converted III into I. II were derived from (2R,3S)-4-oxo-3-phenylacetamidoazetidine-2-sulfinic acids by a novel oxidative desulfinylation induced by Pb(OAc)4. In the second strategy, (1S,5R)-analogs of II were treated with pyruvic acid to give (3S,4S)-3-phenylacetamido-4-pyruvoyloxyazetidin-2-ones. Photolysis with loss of MeSH gave I (R = Me). I (R = CH2OMe) was transformed into I

(R = H) by CF3CO2H. I (R = H) lacked antibacterial activity and β -lactamase-inhibitory properties.

IT 121003-03-0P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation, esterification, and bactericidal activity of)

RN 121003-03-0 HCAPLUS

CN 1-Azetidineacetic acid, α -(1-methylethylidene)-2,4-dioxo-3-

[(phenylacetyl)amino] - (9CI) (CA INDEX NAME)

$$\begin{array}{c} \text{CMe}_2 \\ \parallel \\ \text{C-Co}_2\text{H} \\ \\ \text{Ph-CH}_2\text{-C-NH} \\ \end{array}$$

L15 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1972:112987 HCAPLUS

DOCUMENT NUMBER: 76:112987

TITLE: Synthesis of malonimide derivatives as potential

penicillin analogs

AUTHOR(S): Golik, Uri

CORPORATE SOURCE: Dep. Chem., Weizmann Inst. Sci., Rehovot, Israel SOURCE: Journal of Heterocyclic Chemistry (1972), 9(1), 21-4

CODEN: JHTCAD; ISSN: 0022-152X

DOCUMENT TYPE: Journal LANGUAGE: English

GI For diagram(s), see printed CA Issue.

AB Malonimidoacetic acid derivs. (I, II) were synthesized as potential penicillin analogs, but they failed to inhibit the growth of bacteria when

tested in vitro against a range of gram-pos. and gram-neg. microorganisms.

IT : 35359-55-8P

RN

RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of)

CN 1-Azetidineacetic acid, 3,3-diethyl-2,4-dioxo- (9CI) (CA INDEX NAME)

=> FIL REGISTRY COST IN U.S. DOLLARS

COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION FULL ESTIMATED COST 23.54 781.21

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL

CA SUBSCRIBER PRICE

ENTRY SESSION -1.56 -7.80

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REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

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=>

Uploading C:\Program Files\Stnexp\Queries\10519835h.str



chain nodes:
5 6 7 8 9 10
ring nodes:
1 2 3 4
chain bonds:
1-6 3-5 4-7 7-8 8-9 8-10
ring bonds:
1-2 1-4 2-3 3-4
exact/norm bonds:

1-2 1-4 1-6 2-3 3-4 3-5 4-7 7-8 8-9 8-10 isolated ring systems :

containing 1 :

G1:H,M

G2:A,Cb,Cy,Hy,Ak,Ph

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS 10:CLASS

L16 STRUCTURE UPLOADED

=> d 116 L16 HAS NO ANSWERS L16 STR

G1 H, M

G2 A, Cb, Cy, Hy, Ak, Ph

Structure attributes must be viewed using STN Express query preparation.

=> s 116

GENERIC GROUP NOT VALID HERE

Generic groups may not be used in these circumstances:

- 1. Any generic group node (e.g., Hy) in a ring.
- 2. An Ak node attached to another Ak node.

=> Uploading C:\Program Files\Stnexp\Queries\10519835i.str

التاريخ

chain nodes :

5 6 7 8 9 10 13 15

ring nodes : 1 2 3 4

chain bonds :

1-6 2-15 3-5 4-7 7-8 8-9 8-10 9-13

ring bonds :

1-2 1-4 2-3 3-4

exact/norm bonds :

1-2 1-4 1-6 2-3 2-15 3-4 3-5 4-7 7-8 8-9 8-10 9-13

isolated ring systems :

containing 1 :

G1:H,M

G2:Cb,Cy,Hy,Ak,Ph

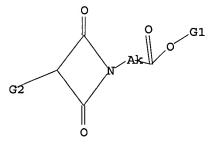
Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS

10:CLASS 13:CLASS 15:CLASS

L17 STRUCTURE UPLOADED

=> d l17 L17 HAS NO ANSWERS L17 STR



G1 H, M

G2 Cb, Cy, Hy, Ak, Ph

Structure attributes must be viewed using STN Express query preparation.

=> s 117

SAMPLE SEARCH INITIATED 14:49:23 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 350 TO ITERATE

,

SEARCH TIME: 00.00.01

100.0% PROCESSED

350 ITERATIONS

0 ANSWERS

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**

PROJECTED ITERATIONS:

5878 TO 8122

05/18/2007

Page 25

PROJECTED ANSWERS: 0 TO 0

0 SEA SSS SAM L17

=> s 117 sss full

FULL SEARCH INITIATED 14:49:30 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED - 6618 TO ITERATE

100.0% PROCESSED 6618 ITERATIONS

SEARCH TIME: 00.00.01

L19 1 SEA SSS FUL L17

=> FIL HCAPLUS

COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION FULL ESTIMATED COST 174.35 955.56

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL ENTRY SESSION CA SUBSCRIBER PRICE 0.00 -7.80

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FILE COVERS 1907 - 18 May 2007 VOL 146 ISS 22 FILE LAST UPDATED: 15 May 2007 (20070515/ED)

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=> s 119

L20 1 L19

=> d 120 ibib abs hitstr tot

L20 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2007 ACS on STN

1972:112987 HCAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 76:112987

TITLE: Synthesis of malonimide derivatives as potential

penicillin analogs

AUTHOR (S): Golik, Uri

CORPORATE SOURCE: Dep. Chem., Weizmann Inst. Sci., Rehovot, Israel SOURCE:

Journal of Heterocyclic Chemistry (1972), 9(1), 21-4

1 ANSWERS

CODEN: JHTCAD; ISSN: 0022-152X

DOCUMENT TYPE: Journal

LANGUAGE: English GT For diagram(s), see printed CA Issue. AB Malonimidoacetic acid derivs. (I, II) were synthesized as potential penicillin analogs, but they failed to inhibit the growth of bacteria when tested in vitro against a range of gram-pos. and gram-neg. microorganisms. 35359-55-8P IT RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of) 35359-55-8 HCAPLUS RN CN-1-Azetidineacetic acid, 3,3-diethyl-2,4-dioxo- (9CI) (CA INDEX NAME) CH2- CO2H Εt Et => s imidoalkan 0 IMIDOALKAN L21 0 IMIDOALKAN => s imidoalkanper 0 IMIDOALKANPER L22 0 IMIDOALKANPER => s imidocarboxylic acids 22 IMIDOCARBOXYLIC 1574229 ACIDS L23 6 IMIDOCARBOXYLIC ACIDS (IMIDOCARBOXYLIC (W) ACIDS) => s 123 and crystalline 77976 CRYSTALLINE 249 CRYSTALLINES 78204 CRYSTALLINE (CRYSTALLINE OR CRYSTALLINES) 356565 CRYST 1801 CRYSTS 357833 CRYST (CRYST OR CRYSTS) 383187 CRYSTALLINE (CRYSTALLINE OR CRYST) L24 0 L23 AND CRYSTALLINE => s 123 and alkanper 0 ALKANPER L25 0 L23 AND ALKANPER => s 123 and alkan 466 ALKAN

05/18/2007

L26

(ALKAN OR ALKANS)

4 ALKANS 470 ALKAN

0 L23 AND ALKAN

```
=> s percarboxylic acids
           278 PERCARBOXYLIC
       1574229 ACIDS
           155 PERCARBOXYLIC ACIDS
L27
                 (PERCARBOXYLIC(W) ACIDS)
=> s 127 and imidoalkan
             0 IMIDOALKAN
L28
             0 L27 AND IMIDOALKAN
=> s 127 and imido
          5172 IMIDO
            14 IMIDOS
          5173 IMIDO
                 (IMIDO OR IMIDOS)
L29
             2 L27 AND IMIDO
=> s 127 and crystalline
         77976 CRYSTALLINE
           249 CRYSTALLINES
         78204 CRYSTALLINE
                 (CRYSTALLINE OR CRYSTALLINES)
        356565 CRYST
          1801 CRYSTS
        357833 CRYST
                 (CRYST OR CRYSTS)
        383187 CRYSTALLINE
                 (CRYSTALLINE OR CRYST)
L30
             0 L27 AND CRYSTALLINE
=> d his
     (FILE 'HOME' ENTERED AT 14:33:37 ON 18 MAY 2007)
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L1
                STRUCTURE UPLOADED
L_2
              0 S L1
L3
              0 S L1 SSS FULL
L4
                STRUCTURE UPLOADED
L5
              1 S L4
L6
              2 S L4 SSS FULL
     FILE 'HCAPLUS' ENTERED AT 14:36:14 ON 18 MAY 2007
L7
              1 S L6
     FILE 'REGISTRY' ENTERED AT 14:38:51 ON 18 MAY 2007
L8
               STRUCTURE UPLOADED
L9
              0 S L8
L10
             11 S L8 SSS FULL
     FILE 'HCAPLUS' ENTERED AT 14:39:18 ON 18 MAY 2007
              7 S L10
L11
     FILE 'REGISTRY' ENTERED AT 14:42:37 ON 18 MAY 2007
L12
               STRUCTURE UPLOADED
L13
              0 S L12
L14
              2 S L12 SSS FULL
     FILE 'HCAPLUS' ENTERED AT 14:43:07 ON 18 MAY 2007
L15
              2 S L14
```

```
FILE 'REGISTRY' ENTERED AT 14:46:07 ON 18 MAY 2007
 L16
               STRUCTURE UPLOADED
 L17
               STRUCTURE UPLOADED
 L18
             0 S L17
 L19
             1 S L17 SSS FULL
     FILE 'HCAPLUS' ENTERED AT 14:49:36 ON 18 MAY 2007
 L20
            1 S L19
 L21
             0 S IMIDOALKAN
 L22
            0 S IMIDOALKANPER
 L23
            6 S IMIDOCARBOXYLIC ACIDS
 L24
            0 S L23 AND CRYSTALLINE
L25
            0 S L23 AND ALKANPER
L26
            0 S L23 AND ALKAN
L27
         155 S PERCARBOXYLIC ACIDS
L28
            0 S L27 AND IMIDOALKAN
L29
             2 S L27 AND IMIDO
 L30
             0 S L27 AND CRYSTALLINE
 => d l29 ibib abs hitstr tot
 L29 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1991:583094 HCAPLUS
 DOCUMENT NUMBER:
                      115:183094
 TITLE:
                      Continuous preparation of imido
                      percarboxylic acids
 INVENTOR(S):
                      Fuchs, Hermann; Gethoeffer, Hanspeter; Gilb, Walter
 PATENT ASSIGNEE(S):
                     Hoechst A.-G., Germany
 SOURCE:
                       Ger. Offen., 4 pp.
                      CODEN: GWXXBX
DOCUMENT TYPE:
                       Patent
LANGUAGE:
                       German
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:
     DE 40000
                                      APPLICATION NO.
                                                            DATE
                             -----
                                        -----
     DE 4003309
                        A1
                             19910808
                                      DE 1990-4003309
                                                             19900205
     EP 441235 A2
                                      EP 1991-101176
                             19910814
                                                             19910130
                             19920226
     EP 441235
                       A3
        R: BE, CH, DE, FR, GB, IT, LI, NL
     US 5132431 A
                          19920721 US 1991-649619
                                                            19910201
                             19910806 CA 1991-2035601
19920803 JP 1991-13473
                    A1
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     JP 04211057
                                                             19910204
 PRIORITY APPLN. INFO.:
                                        DE 1990-4003309 A 19900205
 OTHER SOURCE(S):
                      MARPAT 115:183094
```

GI

AB Title compds. I [n = 2; A = alkylene, alkenylene, phenylene, naphthylene; X = C1-19 alkylene, phenylene] were prepared continuously by oxidizing I (n = 1) in H2SO4 or MeSO3H with organic H2O2 in a static mixer. Thus, \(\varepsilon\)-phthalimidocaproic acid in H2SO4 was treated with 50% aqueous H2O2 to give 91% peracid.

L29 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1990:76941 HCAPLUS

DOCUMENT NUMBER: 112:76941

TITLE: Novel imido aromatic percarboxylic

acids and their preparation and use as

bleaching agents

INVENTOR(S):
Venturello, Carlo; Cavallotti, Caludio

PATENT ASSIGNEE(S): Ausimont S.p.A., Italy SOURCE: Eur. Pat. Appl., 8 pp.

CODEN: EPXXDW

DOCUMENT TYPE:

LANGUAGE:

Patent English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 325288	A1	19890726	EP 1989-101002	19890120
EP 325288	B1	19930901		
R: AT, BE, CH,	DE, ES	, FR, GB, LI,	NL, SE	
US 5575947	Α	19961119	US 1989-299017	19890119
AU 8928679	Α	19890720	AU 1989-28679	19890120
AU 620067	B2	19920213		
JP 02196771	Α	19900803	JP 1989-11743	19890120
JP 2786223	B2	19980813		
AT 93849	T	19930915	AT 1989-101002	19890120
ES 2058347	T 3	19941101	ES 1989-101002	19890120
CA 1340679	C	19990727	CA 1989-588844	19890120
BR 8900264	Α	19890919	BR 1989-264	19890123
PRIORITY APPLN. INFO.:			IT 1988-19132 A	19880120
			EP 1989-101002 A	19890120

OTHER SOURCE(S):

MARPAT 112:76941

GI For diagram(s), see printed CA Issue.

AB Title per acids I [A = (un) substituted benzene or naphthalene nucleus; R = H, CO2H, C(O)OOH, (un) substituted alkyl; n = 1-5] are prepared as bleaching agents. Thus, 44 g 70% H2O2 was added gradually to 55 g phthalimidoacetic acid in 330 g MeSO3H at < 15°, and after 1.5 h the mixture was poured into 20% (NH4)2SO4 at 5°. Filtration, neutralization to pH 6 by Na2CO3 in aqueous Na2SO4, refiltration, aqueous washing, and drying in vacuo at room temperature gave 58 g (97%) of substantially pure phthalimidoperacetic acid

(II). In a bleaching test at 200 mg/L initial active 0, 1.46 g II gave 83.6% bleaching of standard stained cotton, vs. only 75.1% by 1.86 g H-48 (Mg monoperphthalate).

=> log y		
COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	39.53	995.09
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	-2.34	-10.14

STN INTERNATIONAL LOGOFF AT 14:55:55 ON 18 MAY 2007